THURSDAY SEPT 23, 2021

8:45 Welcome and opening remars: Organizing Committee

Interactions between ion transporters and the plasma membrane

9:00-9:30 Gunnar von Heijne, Stockholm University, Sweden: *Biogenesis of integral membrane proteins*

9:30-10:00 Sara Liin, Linköping University, Sweden: Natural and synthetic lipids as activators of voltage-gated potassium channels

10:00-10:15 Oral presentation: Elisa De Grandis Alternating Hemiplegia of Childhood: genotype-phenotype correlations in a cohort of 39 Italian patients

10:15-10:45 Steve Karlish, Weizman institute, Israel: Specific Na,K-ATPase-lipid interactions. A role in neurological disease?

Break 10:45-11:00

11:00-11:30 Erdinc Sezgin, Karolinska Institutet, Sweden: *Structure and function relationship in plasma membrane*

11:30-12:00 Maria Dahlin, Karolinska Institutet/University Hospital, Sweden: *The ketogenic diet - clinical aspects and mechanisms of action*

12:00-12:30 General discussion

Lunch break 12:30-13:15

Pain

13:15-14:00 Håkan Olausson, Linköping university, Sweden: The physiology of pain and novel aspects of pain recording and diagnosis (preliminary title)

14:00-14:30 Philipp Mittermaier, Karolinska University Hospital, Sweden: *Pediatric aspects of pain and pain treatment (preliminary title)*

14:30-15:15 Cecilia Lidbeck and Kickie Löwing, Karolinska Institutet/Karolinska University Hospital, Sweden:

Exploring motor function and everyday activity in a child with ATP1A3

Break 15:15-15:30

15:30-15:45 Oral presentation: Michael Habeck Cryo-EM structures of human Na,K-ATPase α1 and α3 isoforms

15:45-16:00 Oral presentation: Sofia Ygberg Molecular pathology of ATP1A mutations

16:00-16:30 Kathleen Sweadner, Harvard university, USA: The impact of cellular responses on the spectrum of ATP1A3 disease

16:30-17:00 General discussion

FRIDAY SEPT 24, 2021

9:00-10:00 <u>Keynote lecture</u>: Majken Nedergaard, Copenhagen University, Denmark: *The Glymphatic System*

10:00-10:15 Oral presentation: Maria Papadopoulou Sleep disorders in children with Alternating Hemiplegia of Childhood: preliminary results of the HEPNOS study

ATP1A3 in motor neurons, role for motor function

10:15-10:45 Ihtsham Ul Haq, University of Miama, USA: Rapid onset-dystonia-parkinsonism is a movement disorder (preliminary title)

10:45-11:15 Gareth Miles, St Andrews University, UK: *Gait control; role of motor neurons (preliminary title)*

Break 11:15-11:30

11:30-11:45 Oral presentation: To be announced

11:45-12:00 Oral presentation: Frankie Sorell Effects of the Rapid-Onset Dystonia-Parkinsonism gene mutation T613M on spinal motor networks in mice

12:00-12:30 Eva Weidenhielm Broström, Karolinska Institutet/Karolinska University Hospital, Sweden: Gait and motion analysis in pediatric disorders causing motor dysfunction, - how, what and why?

Lunch break 12:30-13:15

Molecular manifestation and genetics

13:15-13:45 Hendrik Rosewich, Göttingen University, Germany: Identifying patients suffering from AHC by screening more children with neurologic symptoms (preliminary title) 13:45-14:15 Anna Lindstrand, Karolinska Institutet, Sweden: Reverse phenotyping after whole genome sequencing allows for high diagnostic rates across a broad spectrum of rare diseases

14:15-14:30 Oral presentation: Monica Dahlstrup Sietam *Spontaneous development of motor deficits in a mouse model carrying the CAPOS mutation*

Break 14:30-14:45

14:45-15:30 Christopher A. Walsh, Harvard Medical School, USA: *Early role for a Na*⁺,*K*⁺-*ATPase (ATP1A3) in brain development*

15:30-16:00 Mohamad Mikati, Duke University, USA: *Gene Therapy of Alternating Hemiplegia of Childhood*

16:00-16:30 General discussion and closing remarks